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=> d his
     (FILE 'HOME' ENTERED AT 09:23:47 ON 02 AUG 2005)
     FILE 'REGISTRY' ENTERED AT 09:23:53 ON 02 AUG 2005
L1
           26021 S EGEG/SQSP
     FILE 'HCAPLUS' ENTERED AT 09:25:27 ON 02 AUG 2005
L2
            4892 S L1
L3
             643 S (FUSION OR CHIMER? OR CHIMAER?) (5A) PROTEIN? AND L2
L4
               6 S L3 AND CAPTUR?
L5
               0 S L3 AND POLYCATION?
               2 S L2 AND POLYCATION?
L6
                 SELECT L6 RN 1-2
L7
          361075 S E1-828
     FILE 'REGISTRY' ENTERED AT 09:32:39 ON 02 AUG 2005
L8
               0 S (AG) \{3-6\} EG \{14-36\} / SOSP
L9
          271929 S (AG) {0-8}EG{2-40}/SOSP
          271887 S .{2-}(AG){0-8}EG{2-40}|(AG){0-8}EG{2-40}.{2-}/SQSP
L10
L11
          271858 S SQL>=6 AND L10
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L12
           32705 S L11
L13
           10387 S L12 AND ?CATION?
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L14
           26011 \text{ S} \cdot \{2-\} \text{ (AG) } \{0-8\} \text{ (EG) } \{2-40\} \text{ (AG) } \{0-8\} \text{ (EG) } \{2-40\} \cdot \{2-\} / \text{SOSP}
L15
             146 S .{2-}(AG)(0-8)(PEG){2-40}|(AG)(0-8)(PEG){2-40}.{2-}/SQSP
L16
           26156 S L14 OR L15
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L17
            4920 S L16
L18
            2014 S L17 AND ?CATION?
L19
              15 S L18 AND CAPTUR?
L20
              25 S L18 AND SOLID(3A) SUPPORT?
L21
               0 S L20 AND TAIL?
L22
              44 S L18 AND TAIL?
L23
               1 S L18 AND ?POLYMER? (3A) TAIL?
L24
               3 S L18 AND TETHER?
L25
            2011 S L18 NOT L24
               2 S L25 AND POSITIV? AND NEGATIV?
L26
L27
            2009 S L25 NOT L26
L28
               1 S L27 AND POLYLYSINE?
L29
               2 S L17 AND POLYLYSINE?
     FILE 'REGISTRY' ENTERED AT 11:17:51 ON 02 AUG 2005
L30
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     FILE 'REGISTRY' ENTERED AT 11:27:21 ON 02 AUG 2005
L31
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L32
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L33
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L34
           26156 S L32 OR L33
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L35

L36

4920 S L34

1 S L27 AND POLYARGININE

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L37
              1 S L17 AND POLYARGININE
L38
              1 S L17 AND POLYARGININE?
L39
              2 S L27 AND ANION? (2A) BIND?
L40
              4 S L27 AND CATION(2A)BIND?
L41
              2 S L27 AND ?ANION?(2A)(BIND? OR BOUND?)
L42
             51 S L27 AND ?CATION? (2A) BIND?
L43
              0 S L27 AND ?CATION? (2A) BOUND?
L44
             1 S L42 AND MATRI?
L45
             50 S L42 NOT L44
L46
             79 S L4 OR L6 OR L19 OR L24 OR L28 OR L29 OR L36-L45
L47
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                DELETE SELECT
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                DELETE SELECT
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     FILE 'HCAPLUS' ENTERED AT 12:11:23 ON 02 AUG 2005
L48
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                                      4216 TERMS
     FILE 'REGISTRY' ENTERED AT 12:11:25 ON 02 AUG 2005
L49
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L50
             58 S L49 AND L16
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L51
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=> d ibib abs 151 1-13
L51 ANSWER 1 OF 13 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER:
                         2003:42910 HCAPLUS
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DOCUMENT NUMBER:

138:102669

TITLE:

Engineered construction of long-wavelength variants of Aequorea green fluorescent protein by computational

modeling from its three-dimensional crystal structure

INVENTOR(S):

Wachter, Rebekka M.; Remington, S. James

PATENT ASSIGNEE(S):

USA

SOURCE:

U.S. Pat. Appl. Publ., 100 pp., Cont.-in-part of U.S.

6.077,707.

CODEN: USXXCO

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND DATE	APPLICATION NO.	DATE
US 2003013149	A1 20030116		20000519
US 6593135 US 6124128	B2 20030715 A 20000926		19960830 <
US 6054321	A 20000425	US 1997-911825	19970815 <
EP 1508574	A2 20050223		19970815
R: AT, BE, CH, IE, FI	DE, DK, ES, FR,	GB, GR, IT, LI, LU, NL,	SE, MC, PT,
US 6077707	A 20000620	US 1997-974737	19971119 <
AU 767375	B2 20031106	AU 2001-23196	20010223
CA 2408302	AA 20011129	CA 2001-2408302	20010517
WO 2001090147	A2 20011129	WO 2001-US16149	20010517
WO 2001090147	A3 20020502		
W: AE, AG, AL,	AM, AT, AU, AZ,	BA, BB, BG, BR, BY, BZ,	CA. CH. CN.

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CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM,
             HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS,
             LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO,
             RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN,
             YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
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     EP 1285065
                          A2
                                20030226
                                            EP 2001-937550
                                                                    20010517
             AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
     JP 2004502410
                          T2
                                20040129
                                             JP 2001-586334
                                                                    20010517
     US 2004014128
                                20040122
                          A1
                                             US 2003-620099
                                                                    20030714
PRIORITY APPLN. INFO.:
                                             US 1996-24050P
                                                                 P 19960816
                                             US 1996-706408
                                                                 A1 19960830
                                             US 1997-911825
                                                                 A1 19970815
                                             US 1997-974737
                                                                 A2 19971119
                                            AU 1997-43277
                                                                 A3 19970815
                                                                 A3 19970815
                                            'EP 1997-941350
                                             US 2000-575847
                                                                 A 20000519
                                             WO 2001-US16149
                                                                 W .20010517
AB
     Engineered fluorescent proteins, nucleic acids encoding them, and methods
     of use are provided. As a step in understanding the properties of green
     fluorescent protein (GFP) from Aequorea victoria, and to aid in the
     tailoring of GFPs with altered characteristics, the 3-dimensional
     structure was determined at 1.9 Å resolution of the S65T mutant. Spectral
     properties of Thr203 mutants in comparison to S65T are provided.
     particular, the S65G/V68L/S72A/T203Y (designated yellow fluorescent
     protein or YFP) displays an excitation maximum at 514 nm, an emission maximum
at.
     527 nm, extinction coefficient of 83,400 M-1cm-1, and quantum yield of 0.61;
     its absorption spectrum is a function of NaCl concentration, demonstrating its
     usefulness as a halide sensor. Crystallog. identification and
     description of halide binding sites indicates a relationship between
     anion binding and cavity size, a relaxation of the
     \beta-barrel conformation in response to the H148Q substitution and
     iodide binding, and the key residues for anion binding
     are determined by mutational anal.
    ANSWER 2 OF 13 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER:
                         2000:717215 HCAPLUS
DOCUMENT NUMBER:
                         134:176440
TITLE:
                         Identification of nucleolar protein No55 as
                         a tumour-associated autoantigen in patients with
                         prostate cancer
AUTHOR(S):
                         Fossa, A.; Siebert, R.; Aasheim, H.-C.; Maelandsmo, G.
                         M.; Berner, A.; Fossa, S. D.; Paus, E.; Smeland, E.
                         B.; Gaudernack, G.
CORPORATE SOURCE:
                         Department of Immunology, The Norwegian Radium
                         Hospital, Oslo, 0310, Norway
SOURCE:
                         British Journal of Cancer (2000), 83(6),
                         743-749
                         CODEN: BJCAAI; ISSN: 0007-0920
PUBLISHER:
                         Harcourt Publishers Ltd.
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     Four different genes were identified by immunoscreening of a cDNA
     expression library from the human prostate cancer cell line DU145 with
     allogeneic sera from four prostate cancer patients. A cDNA encoding the
```

nucleolar protein No55 was further analyzed and shown to be expressed at

the mRNA level in several normal tissues, including ovaries, pancreas and prostate and in human prostate cancer cell lines PC-3, PC-3m and LNCaP. By reverse transcriptase/polymerase chain reaction, expression of No55 was several-fold higher in two out of nine prostate cancer primary tumors and two out of two metastatic lesions, compared to normal prostate tissue. Antibodies to No55 were detected in sera from seven out of 47 prostate cancer patients but not in sera from 20 healthy male controls. Sequence anal. of the No55 open reading frame from normal and tumor tissues revealed no tumor-specific mutations. The No55 gene was located to chromosome 17q21, a region reported to be partially deleted in prostate cancer. Considering the immunogenicity of the No55 protein in the tumor host, the expression profile and chromosomal localization of the corresponding gene, studies evaluating No55 as a potential antigen for immunol. studies in prostate cancer may be warranted.

REFERENCE COUNT: 32 THERE ARE 32 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L51 ANSWER 3 OF 13 HCAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2000:697875 HCAPLUS

DOCUMENT NUMBER: 134:290979

TITLE: Isoforms of JSAP1 scaffold protein generated through

alternative splicing

AUTHOR(S): Ito, M.; Akechi, M.; Hirose, R.; Ichimura, M.;

Takamatsu, N.; Xu, P.; Nakabeppu, Y.; Tadayoshi, S.;

Yamamoto, K.-i.; Yoshioka, K.

CORPORATE SOURCE: School of Science, Department of Biosciences, Kitasato

University, Sagamihara, Kanagawa, 228-8555, Japan

SOURCE: Gene (2000), 255(2), 229-234

CODEN: GENED6; ISSN: 0378-1119

PUBLISHER: Elsevier Science B.V.

DOCUMENT TYPE: Journal LANGUAGE: English

We have identified four isoforms of c-Jun NH2-terminal kinase (JNK)/stress-activated protein kinase-associated protein 1 (JSAP1), a scaffold protein that participates in JNK mitogen-activated protein kinase cascades, termed JSAPla, JSAPlb, JSAPlc, and JSAPld. The previously identified JSAP1 was renamed JSAP1a to avoid confusion. Analyses of the exon-intron structure of the jsapl gene indicated that the isoforms are generated through alternative splicing involving exons 5 and 6. The mRNA expression levels of the JSAP1 isoforms differed among the mouse tissues examined We also investigated the region of JSAP1 responsible for its interaction with JNK, and found that the JNK-binding domain is located between aa residues 201 and 217 in JSAPla, which is encoded by part of exon 6. As all the JSAP1 isoforms contain this binding domain, we examined the binding affinity of the JSAP1 isoforms for JNK1, JNK2, and JNK3. JSAP1c and JSAP1d, which contain a 31-aa sequence not present in JSAP1a or JSAP1b, had a lower binding affinity for the JNKs, especially JNK3. These results suggest that JSAP1c and JSAP1d may attenuate the scaffolding activity of JSAPla and/or JSAPlb in JNK cascades, especially the JNK3 cascades.

REFERENCE COUNT: 20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L51 ANSWER 4 OF 13 HCAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2000:627482 HCAPLUS

DOCUMENT NUMBER: 133:262900

TITLE: Sequence similarities between a novel putative G

protein-coupled receptor and Na+/Ca2+ exchangers

define a cation binding domain

AUTHOR(S): Nikkila, Heli; McMillan, D. Randy; Nunez, Brian S.;

Pascoe, Leigh; Curnow, Kathleen M.; White, Perrin C.

CORPORATE SOURCE: Division of Pediatric Endocrinology, University of

Texas Southwestern Medical Center, Dallas, TX,

75235-9063, USA

SOURCE: Molecular Endocrinology (2000), 14(9),

1351-1364

CODEN: MOENEN; ISSN: 0888-8809

PUBLISHER: Endocrine Society

DOCUMENT TYPE: Journal LANGUAGE: English

CDNA clones encoding a novel putative G protein-coupled receptor have been characterized. The receptor is widely expressed in normal solid tissues. Consisting of 1967 amino acid residues, this receptor is one of the largest known and is therefore referred to as a very large G protein-coupled receptor, or VLGR1. It is most closely related to the secretin family of G protein-coupled receptors based on similarity of the sequences of its transmembrane segments. As demonstrated by cell surface labeling with a biotin derivative, the recombinant protein is expressed on the surface of transfected mammalian cells. Whereas several other recently described receptors in this family also have large extracellular domains, the large extracellular domain of VLGR1 has a unique structure. It has nine imperfectly repeated units that are rich in acidic residues and are spaced at intervals of approx. 120 amino acid residues. These repeats resemble the regulatory domains of Na+/Ca2+ exchangers as well as a component of an extracellular aggregation factor of marine sponges. Bacterial fusion proteins containing two or four repeats specifically bind 45Ca in overlay expts.; binding is competed poorly by Mg2+ but competed well by neomycin, Al3+, and Gd3+. These results define a consensus cation binding motif employed in several widely

divergent types of proteins. The ligand for VLGR1, its function, and the signaling pathway(s) it employs remain to be defined.

REFERENCE COUNT:

THERE ARE 41 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L51 ANSWER 5 OF 13 HCAPLUS COPYRIGHT 2005 ACS on STN

41

ACCESSION NUMBER: 2000:613167 HCAPLUS

DOCUMENT NUMBER: 133:218310

TITLE: DNA sequence of both chromosomes of the cholera

pathogen Vibrio cholerae

AUTHOR(S): Heidelberg, John F.; Elsen, Jonathan A.; Nelson,

William C.; Clayton, Rebecca A.; Gwinn, Michelle L.; Dodson, Robert J.; Haft, Daniel H.; Hickey, Erin K.; Peterson, Jeremy D.; Umayam, Lowell; Gill, Steven R.; Nelson, Karen E.; Read, Timothy D.; Tettelin, Herve; Richardson, Delwood; Ermolaeva, Maria D.; Vamathevan, Jessica; Bass, Steven; Qin, Haiying; Dragoi, Loana; Sellers, Patrick; McDonald, Lisa; Utterback, Teresa; Fleishmann, Robert D.; Nierman, William C.; White, Owen; Salzberg, Steven L.; Smith, Hamilton O.;

Colwell, Rita R.; Mekalanos, John J.; Venter, J. Craig; Fraser, Claire M.

CORPORATE SOURCE: The Institute for Genomic Research, Rockville, MD,

20850, USA

SOURCE: Nature (London) (2000), 406(6795), 477-483

CODEN: NATUAS; ISSN: 0028-0836

PUBLISHER: Nature Publishing Group

DOCUMENT TYPE: Journal LANGUAGE: English

AB The complete genomic sequence of the gram-neg., γ -Proteobacterium Vibrio cholerae El Tor N16961 was determined to be 4,033,460 bp. The genome consists of two circular chromosomes of 2,961,146 bp and 1,072,314 bp that

together encode 3885 open reading frames. The vast majority of recognizable genes for essential cell functions (such as DNA replication, transcription, translation, and cell-wall biosynthesis) and pathogenicity (for example, toxins, surface antigens, and adhesins) are located on the large chromosome. In contrast, the small chromosome contains a larger fraction (59%) of hypothetical genes compared with the large chromosome (42%), and also contains many more genes that appear to have origins other than the $\gamma\textsc{-Proteobacteria}$. The small chromosome also carries a gene capture system (the integron island) and host 'addiction' genes that are typically found on plasmids; thus, the small chromosome may have originally been a megaplasmid that was captured by an ancestral Vibrio species. The V. cholerae genomic sequence provides a starting point for understanding how a free-living, environmental organism emerged to become a significant human bacterial pathogen.

REFERENCE COUNT: 50 THERE ARE 50 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L51 ANSWER 6 OF 13 HCAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2000:4884 HCAPLUS

DOCUMENT NUMBER: 132:163331

TITLE: env sequences of simian immunodeficiency viruses from

chimpanzees in Cameroon are strongly related to those of human immunodeficiency virus group N from the same

geographic area

AUTHOR(S): Corbet, Sylvie; Muller-Trutwin, Michaela C.;

Versmisse, Pierre; Delarue, Severine; Ayouba, Ahidjo;

Lewis, John; Brunak, Soren; Martin, Paul; Brun-Vezinet, Francoise; Simon, Francois; Barre-Sinoussi, Francoise; Mauclere, Philippe

CORPORATE SOURCE: Unite de Biologie des Retrovirus, Institut Pasteur,

Paris, Fr.

SOURCE: Journal of Virology (2000), 74(1), 529-534

CODEN: JOVIAM; ISSN: 0022-538X

PUBLISHER: American Society for Microbiology

DOCUMENT TYPE: Journal LANGUAGE: English

Human immunodeficiency virus type 1 (HIV-1) group N from Cameroon is phylogenetically close, in env, to the simian immunodeficiency virus (SIV) cpz-qab from Gabon and SIVcpz-US of unknown geog. origin. We screened 29 wild-born Cameroonian chimpanzees and found that three (Cam3, Cam4, and Cam5) were pos. for HIV-1 by Western blotting. Mitochondrial DNA sequence anal. demonstrated that Cam3 and Cam5 belonged to Pan troglodytes troglodytes and that Cam4 belonged to P. t. vellerosus. Genetic analyses of the viruses together with serol. data demonstrated that at least one of the two P. t. troglodytes chimpanzees (Cam5) was infected in the wild, and revealed a horizontal transmission between Cam3 and Cam4. These data confirm that P. t. troglodytes is a natural host for HIV-1-related viruses. Furthermore, they show that SIVcpz can be transmitted in captivity, from one chimpanzee subspecies to another. All three SIVcpz-cam viruses clustered with HIV-1 N in env. The full Cam3 SIVcpz genome sequence showed a very close phylogenetic relationship with SIVcpz-US, a virus identified in a P. t. troglodytes chimpanzee captured nearly 40 yr earlier. Like SIVcpz-US, SIVcpz-cam3 was closely related to HIV-1 N in env, but not in pol, supporting the hypothesis that HIV-1 N results from a recombination event. SIVcpz from chimpanzees born in the wild in Cameroon are thus strongly related in env to HIV-1 N from Cameroon, demonstrating the geog. coincidence of these human and simian viruses and providing a further strong argument in favor of the origin of HIV-1 being in chimpanzees.

REFERENCE COUNT: 27 THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L51 ANSWER 7 OF 13 HCAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1999:766689 HCAPLUS

DOCUMENT NUMBER: 132:133025

TITLE: Identification of a Novel Homolog of the Drosophila staufen Protein in the Chromosome

8q13-q21.1 Region

AUTHOR(S): Buchner, Georg; Bassi, Maria Teresa; Andolfi, Grazia;

Ballabio, Andrea; Franco, Brunella

CORPORATE SOURCE: Telethon Institute of Genetics and Medicine (TIGEM),

Milan, Italy

SOURCE: Genomics (1999), 62(1), 113-118

CODEN: GNMCEP; ISSN: 0888-7543

PUBLISHER: Academic Press

DOCUMENT TYPE: Journal LANGUAGE: English

AB We report the identification of a new transcript homologous to the Drosophila staufen protein. This transcript, named STAU2 (HGMW-approved gene symbol and name), maps to the chromosome 8q13-q21 region. The full-length STAU2 cDNA is 4058 bp and contains an open reading frame of 479 amino acids. Anal. of the predicted protein product indicated the presence of three double-stranded RNA-binding domains. Best-fit anal. revealed a 48.5% similarity to the Drosophila protein and a 59.9% similarity to the recently described mammalian homolog hStau, indicating that at least two different transcripts with homologies to the fly protein are present in mammals. (c) 1999 Academic Press.

REFERENCE COUNT: 15 THERE ARE 15 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L51 ANSWER 8 OF 13 HCAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1999:594265 HCAPLUS

DOCUMENT NUMBER: 131:320676

TITLE: mAKAP: an A-kinase anchoring protein targeted to the

nuclear membrane of differentiated myocytes

AUTHOR(S): Kapiloff, Michael S.; Schillace, Robynn V.; Westphal,

Ann M.; Scott, John D.

CORPORATE SOURCE: Howard Hughes Medical Institute, Vollum Institute,

Portland, OR, 97201-3098, USA

SOURCE: Journal of Cell Science (1999), 112(16),

2725-2736

CODEN: JNCSAI; ISSN: 0021-9533

PUBLISHER: Company of Biologists Ltd.

DOCUMENT TYPE: Journal LANGUAGE: English

The compartmentalization of second messenger-activated protein kinases contributes to the fidelity of hormone-mediated signal transduction events. For example, the cAMP-dependent protein kinase is tethered at specific intracellular locations through association with A-kinase anchoring proteins (AKAPs). The authors now report the cloning of mAKAP, an anchoring protein found predominantly in heart, skeletal muscle and brain, and whose expression is induced in neonatal ventriculocytes by treatment with hypertrophic stimuli. MAKAP is targeted to the nuclear membrane of differentiated myocytes. Anal. of mAKAP-green fluorescent protein (GFP) fusion constructs revealed that nuclear membrane targeting is conferred by two regions of the protein, between residues

772-915 and 915-1065, which contain spectrin-like repeat sequences. Heterologous expression of the mAKAP targeting sequences displaced the endogenous anchoring protein from the nuclear membrane, demonstrating that

mAKAP targeting is saturable. Collectively, these data suggest that a domain containing spectrin-like repeats mediates targeting of the anchoring protein mAKAP and the cAMP-dependent protein kinase holoenzyme to the

nuclear membrane in response to differentiation signals.

REFERENCE COUNT: 41 THERE ARE 41 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L51 ANSWER 9 OF 13 HCAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1998:460763 HCAPLUS

DOCUMENT NUMBER: 129:185689

TITLE: Identification of a human PTS1 receptor

docking protein directly required for peroxisomal

protein import

AUTHOR(S): Fransen, Marc; Terlecky, Stanley R.; Subramani, Suresh

CORPORATE SOURCE: Department of Biology, University of California at San

Diego, La Jolla, CA, 92093-0322, USA

SOURCE: Proceedings of the National Academy of Sciences of the

United States of America (1998), 95(14),

8087-8092

CODEN: PNASA6; ISSN: 0027-8424 National Academy of Sciences

DOCUMENT TYPE: Journal LANGUAGE: English

PUBLISHER:

The discovery of many fatal human disorders resulting from impaired peroxisomal protein import makes the functional characterization of human peroxins critical As part of our attempt to identify novel human genes and gene products involved in the import of peroxisomal proteins, the authors raised antisera against peroxisomal membrane proteins. One such antiserum inhibited peroxisomal protein import in semipermeabilized mammalian cells. This "import inhibiting" antiserum, ab-MF3, specifically recognized a 57-kDa protein. Immunoblot anal. of rat liver subcellular fractions demonstrated that this protein was present exclusively in peroxisomal membranes. Functional anal. revealed that this 57-kDa mol. bound the PTS1 receptor, Tcx5p, in ligand blots, suggesting it is a docking site on the peroxisomal membrane. Previous studies have identified two yeast proteins, Pex14p and Pex13p, as Pex5p-binding proteins. To facilitate the biochem. anal. of peroxisomal membrane docking proteins, the authors cloned and expressed the previously unidentified human Pex14p, as well as a human Pex13p that is 39 aa longer than previously reported. Recombinant Pex14p was specifically recognized by the "import inhibiting" ab-MF3 and bound Pex5p and the Src homol. 3 (SH3) domain of Pex13p in ligand blots. These studies demonstrate that the ab-MF3-immunoreactive, 57-kDa peroxisomal membrane protein is Pex14p. Furthermore, this peroxin interacts with Pex5p and Pex13p(SH3) and is directly required for peroxisomal protein import.

REFERENCE COUNT: 33 THERE ARE 33 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L51 ANSWER 10 OF 13 HCAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1996:424017 HCAPLUS

DOCUMENT NUMBER: 125:110877

TITLE: Association of human protein-tyrosine phosphatase

κ with members of the Armadillo family

AUTHOR(S): Fuchs, Miriam; Mueller, Thomas; Lerch, Markus M.;

Ullrich, Axel

CORPORATE SOURCE: Dep. Molecular Biol., Max-Planck-Inst. Biochemie,

Martinsried, 82152, Germany

SOURCE: Journal of Biological Chemistry (1996),

271(28), 16712-16719

CODEN: JBCHA3; ISSN: 0021-9258

PUBLISHER: American Society for Biochemistry and Molecular

Biology

DOCUMENT TYPE: Journal LANGUAGE: English

AB A human receptor-like protein-tyrosine phosphatase (PTP) was identified in the mammary carcinoma cell line SK-BR-3, which represents the human homolog of murine PTPk and was therefore termed hPTPk. The

hPTP κ expression is dependent on cell d. and is colocalized with 2

members of the arm family of proteins, β -catenin and

 $\gamma\text{-catenin/plakoglobin,}$ at adherens junctions. Both in vitro and in vivo binding assays demonstrated specific complex formation between

endogenous hPTP κ and β - and γ -catenin/plakoglobin. In

addition, evidence that suggests that $\beta\text{-catenin}$ may represent a substrate for the catalytic activity of $\text{hPTP}\kappa$. The

identification of specific binding partners for this

receptor-like PTP provides insight into the mechanisms of its biol. action

and suggests a role for hPTP κ in the regulation of processes involving cell contact and adhesion such as growth control, tumor invasion, and metastasis.

L51 ANSWER 11 OF 13 HCAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1996:137493 HCAPLUS

DOCUMENT NUMBER: 124:222269

TITLE: Identification and characterization of a

thermostable MutS homolog from Thermus aquaticus

AUTHOR(S): Biswas, Indranil; Hsieh, Peggy

CORPORATE SOURCE: Genetics & Biochemistry Branch, NIDDK, Bethesda, MD,

20892-1810, USA

SOURCE: Journal of Biological Chemistry (1996),

271(9), 5040-8

CODEN: JBCHA3; ISSN: 0021-9258

PUBLISHER: American Society for Biochemistry and Molecular

Biology

DOCUMENT TYPE: Journal LANGUAGE: English

Recognition of mispaired or unpaired bases during DNA mismatch repair is AB carried out by the MutS protein family. Here, the authors describe the isolation and characterization of a thermostable MutS homolog from Thermus aquaticus YT-1. Sequencing of the mutS gene predicts an 89.3-kDa polypeptide sharing extensive amino acid sequence homol. with MutS homologs from both prokaryotes and eukaryotes. Expression of the T. aquaticus mutS gene in Escherichia coli results in a dominant mutator phenotype. Initial biochem. characterization of the thermostable MutS protein, which was purified to apparent homogeneity, reveals two thermostable activities, an ATP hydrolysis activity in which ATP is hydrolyzed to ADP and Pi and a specific DNA mismatch binding activity with affinities for heteroduplex DNAs containing either an insertion/deletion of one base or a GT mismatch. The ATPase activity exhibits a temperature optimum of approx. 80°C. Heteroduplex DNA binding by the T. aquaticus MutS protein requires Mg2+ and occurs over a broad temperature range from 0°C to at least 70°C. The thermostable MutS protein may be useful for further biochem. and structural studies of mismatch binding and for applications involving mutation detection.

L51 ANSWER 12 OF 13 HCAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER:

1995:301696 HCAPLUS

DOCUMENT NUMBER:

122:179969

TITLE:

Nucleotide sequence and mutational analysis indicate that two Helicobacter pylori genes encode a P-type

ATPase and a cation-binding

protein associated with copper transport

AUTHOR(S): Ge, Zhongming; Hiratsuka, Koji; Taylor, Diane E. CORPORATE SOURCE:

Dep. Med. Microbiol. Infectious Diseases, Univ.

Alberta, Edmonton, AB, T6G 2H7, Can.

SOURCE: Molecular Microbiology (1995), 15(1), 97-106

CODEN: MOMIEE; ISSN: 0950-382X

PUBLISHER: Blackwell DOCUMENT TYPE: Journal LANGUAGE: English

A 2.7-kb fragment of Helicobacter pylori UA802 chromosomal DNA was cloned and sequenced. Three open reading frames (designated ORF1, ORF2 and ORF3, resp.) were predicted from the DNA sequence, of which ORF1 and ORF2 appeared to be located within the same operon. The deduced 611-amino-acid sequence of ORF1, a P-type ATPase (designated hpCopA), had striking homol. (29-38%) with several bacterial P-type ATPases and contained the potential functional domains conserved in P-type ATPases from various sources ranging from bacterial to human. A protein of 66 amino acids (designated hpCopP) encoded by ORF2 shared extensive sequence similarity with MerP, a periplasmic mercuric ion-transporting protein, and contains the heavy metal-binding motif. Disruption of ORF1 with a chloramphenicol-resistance cassette (CAT) rendered the H. pylori mutants more susceptible to cupric ion than their parental strains, whereas there is no significant alternation of susceptibility to Ni2+, Cd2+ and Hq2+ between the mutants and the parental strains. The results obtained indicate that ORF1 and ORF2 comprise a cation-transporting system which is associated with copper export out of the H. pylori cells.

L51 ANSWER 13 OF 13 HCAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER:

1993:443606 HCAPLUS

DOCUMENT NUMBER:

119:43606

TITLE:

Identification and preliminary

characterization of a protein motif related to the

zinc finger

AUTHOR(S):

Lovering, Ruth; Hanson, Isabel M.; Borden, Katherine L. B.; Martin, Stephen; O'Reilly, Nicola J.; Evan, Gerard I.; Rahman, Dinah; Pappin, Darryl J. C.;

Trowsdale, John; Freemont, Paul S.

CORPORATE SOURCE:

Hum. Immunogenet. Lab., Imp. Cancer Res. Fund, London,

WC2A 3PX, UK

SOURCE:

Proceedings of the National Academy of Sciences of the

United States of America (1993), 90(6),

2112-16

CODEN: PNASA6; ISSN: 0027-8424

DOCUMENT TYPE: LANGUAGE:

Journal English

A protein motif related to the zinc finger was identified which defines a newly discovered family of proteins. The motif was found in the sequence of the human RING1 gene, which is proximal to the major histocompatibility complex region on chromosome 6. The name RING finger is proposed for this motif, which is found in 27 proteins, all of which have putative DNA binding functions. A peptide corresponding to the RING1 motif was synthesized and a number of its properties, including metal and DNA binding, were examined The RING finger motif appears to be the DNA-binding domain of this newly defined family of proteins.

=> d his full

(FILE 'HOME' ENTERED AT 09:23:47 ON 02 AUG 2005)

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L3
L4
              6 SEA L3 AND CAPTUR?
L5
              O SEA L3 AND POLYCATION?
              2 SEA L2 AND POLYCATION?
L6
                D SCAN
                SELECT L6 RN 1-2
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L10
         271887 SEA .{2-}(AG){0-8}EG{2-40}|(AG){0-8}EG{2-40}.{2-}/SQSP
L11
         271858 SEA SQL>=6 AND L10
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L12
          32705 SEA L11
L13
          10387 SEA L12 AND ?CATION?
     FILE 'REGISTRY' ENTERED AT 10:01:10 ON 02 AUG 2005
L14
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L15
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L16
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            2014 SEA L17 AND ?CATION?
L18
L19
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L20
              25 SEA L18 AND SOLID(3A) SUPPORT?
               O SEA L20 AND TAIL?
L21
L22
              44 SEA L18 AND TAIL?
L23
               1 SEA L18 AND ?POLYMER? (3A) TAIL?
                 D SCAN
L24
               3 SEA L18 AND TETHER?
                 D SCAN
L25
            2011 SEA L18 NOT L24
               2 SEA L25 AND POSITIV? AND NEGATIV?
L26
                 D SCAN
. L*** DEL
               0 S L25 NOT L25
L27
            2009 SEA L25 NOT L26
L*** DEL
               0 S BLOCK (A) HOMOPOLYMER? (5A) AMINO (3A) ACID
L*** DEL
              18 S BLOCK(A) POLYMER? (5A) AMINO (3A) ACID
                 D L27 TI 1-20
L28
               1 SEA L27 AND POLYLYSINE?
                 D SCAN
L29
               2 SEA L17 AND POLYLYSINE?
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L30
               1 SEA 775512-26-0
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     FILE 'HCAPLUS' ENTERED AT 11:25:29 ON 02 AUG 2005
                 D SCAN L29
     FILE 'REGISTRY' ENTERED AT 11:27:21 ON 02 AUG 2005
L31
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                 D SQIDE L31
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L32
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L36
               1 SEA L27 AND POLYARGININE
L37
               1 SEA L17 AND POLYARGININE
L38
               1 SEA L17 AND POLYARGININE?
L*** DEL
             624 S L27 AND BIND?
L39
               2 SEA L27 AND ANION? (2A) BIND?
L40
               4 SEA L27 AND CATION(2A)BIND?
                 D SCAN L39
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L*** DEL
               0 S L27 AND POLYANION? (2A) BIND?
L41
               2 SEA L27 AND ?ANION? (2A) (BIND? OR BOUND?)
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L42
              51 SEA L27 AND ?CATION?(2A)BIND?
L43
               O SEA L27 AND ?CATION? (2A) BOUND?
L44
               1 SEA L42 AND MATRI?
                 D SCAN
L45
              50 SEA L42 NOT L44
                 D TI 1-2
                 D TI 2-20
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L46

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L47

13 SEA PY<2001 AND L46

D SCAN

DELETE SELECT

SELECT L47 RN 1-13

DELETE SELECT

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FILE 'HCAPLUS' ENTERED AT 12:11:23 ON 02 AUG 2005. L48 TRA L47 1-13 RN : 4216 TERMS

FILE 'REGISTRY' ENTERED AT 12:11:25 ON 02 AUG 2005

L49 4216 SEA L48

L50 58 SEA L49 AND L16

L*** DEL O S HCAPLUS

FILE 'HCAPLUS' ENTERED AT 12:13:54 ON 02 AUG 2005 L51 13 SEA L47 AND L50 D IBIB ABS L51 1-13

FILE HOME

FILE REGISTRY

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 1 AUG 2005 HIGHEST RN 857935-17-2 DICTIONARY FILE UPDATES: 1 AUG 2005 HIGHEST RN 857935-17-2

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* The CA roles and document type information have been removed from * the IDE default display format and the ED field has been added, * effective March 20, 2005. A new display format, IDERL, is now * available and contains the CA role and document type information. *

Structure search iteration limits have been increased. See HELP SLIMITS for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at: http://www.cas.org/ONLINE/DBSS/registryss.html

FILE HCAPLUS

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Welcome to STN International! Enter x:x
LOGINID: ssspat01dxs
PASSWORD:
 * * * * * * RECONNECTED TO STN INTERNATIONAL * * * * * * S
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     FILE 'REGISTRY' ENTERED AT 09:23:53 ON 02 AUG 2005
L1
          26021 S EGEG/SOSP
     FILE 'HCAPLUS' ENTERED AT 09:25:27 ON 02 AUG 2005
L2
           4892 S L1
L3
            643 S (FUSION OR CHIMER? OR CHIMAER?) (5A) PROTEIN? AND L2
L4
              6 S L3 AND CAPTUR?
L5
              0 S L3 AND POLYCATION?
              2 S L2 AND POLYCATION?
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L9
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L10
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L11
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L12
          32705 S L11
L13
          10387 S L12 AND ?CATION?
     FILE 'REGISTRY' ENTERED AT 10:01:10 ON 02 AUG 2005
L14
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L16
          26156 S L14 OR L15
     FILE 'HCAPLUS' ENTERED AT 10:06:00 ON 02 AUG 2005
L17
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L18
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L19
             15 S L18 AND CAPTUR?
L20
             25 S L18 AND SOLID(3A) SUPPORT?
L21
              0 S L20 AND TAIL?
L22
             44 S L18 AND TAIL?
L23
              1 S L18 AND ?POLYMER? (3A) TAIL?
L24
              3 S L18 AND TETHER?
L25
           2011 S L18 NOT L24
L26
              2 S L25 AND POSITIV? AND NEGATIV?
L27
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FILE 'REGISTRY' ENTERED AT 11:17:51 ON 02 AUG 2005

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· L31
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      L35
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                                                                            1 S L27 AND POLYARGININE
      L37
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      L38
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      L39
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      L40
                                                                            4 S L27 AND CATION(2A)BIND?
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      L41
                                                                        51 S L27 AND ?CATION?(2A)BIND?
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      L43
                                                                           1 S L42 AND MATRI?
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      L45
                                                                       79 S L4 OR L6 OR L19 OR L24 OR L28 OR L29 OR L36-L45
      L46
       L47
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                                                                                       TRA L47 1-13 RN :
                                                                                                                                                                                                    4216 TERMS
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       L49
                                                               4216 SEA L48
       L50
                                                                        58 S L49 AND L16
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      L51
                                                                       13 S L47 AND L50
      L52
                                                                    846 S L17 AND (FUSION OR CHIMER? OR CHIMAER?)
      L53
                                                                   302 S L52 AND BIND?
      L54
                                                                         27 S L53 AND (IMMOBILIZ? OR IMMOBILIS?)
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                                                                         69 S E1-E69
                                                                            3 S L56 AND L16
      L57
      => fil hcaplus
       FILE 'HCAPLUS' ENTERED AT 12:56:27 ON 02 AUG 2005
      USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
       PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s 157

L58 · 4 L57

=> s 157 and 155 4 L57

L59 3 L57 AND L55

=> d sqide 157 1-3

L57 ANSWER 1 OF 3 REGISTRY COPYRIGHT 2005 ACS on STN

RN **283617-63-0** REGISTRY

CN Peptide (synthetic) fusion protein with green fluorescent protein (Aequorea victoria) fusion protein with peptide (synthetic linker) fusion protein with anti-(human chorionic gonadotropin α-subunit) immunoglobulin (Lama glama heavy chain V-D-J region) fusion protein with anti-(red reactive 6) immunoglobulin (Lama glama heavy chain V-D-J region) fusion protein with peptide (synthetic myc tag) (9CI) (CA INDEX NAME) OTHER NAMES:

CN 44: PN: WO0040968 SEQID: 43 claimed protein

FS PROTEIN SEQUENCE

SQL 500

PATENT ANNOTATIONS (PNTE):

Sequence | Patent

Source | Reference

Not Given|WO2000040968

|claimed |SEOID 43

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SR
   CA
LC
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   STN Files:
DT.CA CAplus document type: Patent
     Roles from patents: BIOL (Biological study); PRP (Properties); USES
RL.P
     (Uses)
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           1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
T.57
   ANSWER 2 OF 3 REGISTRY COPYRIGHT 2005 ACS on STN
   158935-45-6 REGISTRY
RN
CN
   5-13-Dynorphin B [5-methionine, 6-alanine] (synthetic clone pLM138) fusion
   protein with protein (synthetic 84-amino acid fragment) (9CI) (CA INDEX
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SQL
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   CA
LC
   STN Files: CA, CAPLUS
DT.CA CAplus document type: Journal
RL.NP Roles from non-patents: PRP (Properties)
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           1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
   ANSWER 3 OF 3 REGISTRY COPYRIGHT 2005 ACS on STN
L57
   138361-59-8 REGISTRY
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CN
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    (9CI) (CA INDEX NAME)
OTHER NAMES:
CN
   Laminin (human A-subunit precursor)
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        151 YYAVSDSECL SRYNITPRRG PPTYRADDEV ICTSYYSRLV PLEHGEIHTS
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LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL DT.CA CAplus document type: Journal; Patent

RL.P Roles from patents: BIOL (Biological study); PROC (Process); PRP

(Properties); USES (Uses) Roles from non-patents: PRP (Properties) RL.NP 2 REFERENCES IN FILE CA (1907 TO DATE) 2 REFERENCES IN FILE CAPLUS (1907 TO DATE) => d 150 sqide 1-58 YOU HAVE REQUESTED DATA FROM FILE 'REGISTRY' - CONTINUE? (Y)/N:v T.50 ANSWER 1 OF 58 REGISTRY COPYRIGHT 2005 ACS on STN 482390-90-9 REGISTRY RN CN Green fluorescent protein [65-glycine,72-alanine,203-tyrosine] (Aequorea victoria) (9CI) (CA INDEX NAME) FS PROTEIN SEQUENCE SQL 238 SEO 1 MSKGEELFTG VVPILVELDG DVNGHKFSVS GEGEGDATYG KLTLKFICTT 51 GKLPVPWPTL VTTFGYGVOC FARYPDHMKR HDFFKSAMPE GYVOERTIFF 101 KDDGNYKTRA EVKFEGDTLV NRIELKGIDF KEDGNILGHK LEYNYNSHNV 151 YIMADKQKNG IKVNFKIRHN IEDGSVQLAD HYQQNTPIGD GPVLLPDNHY 201 LSYOSALSKD PNEKRDHMVL LEFVTAAGIT HGMDELYK HITS AT: 1-238 **RELATED SEQUENCES AVAILABLE WITH SEQLINK** Unspecified CIMAN SR CA STN Files: CA, CAPLUS, TOXCENTER, USPAT2, USPATFULL LC DT.CA CAplus document type: Patent Roles from patents: ANST (Analytical study); BIOL (Biological study); RL.P PREP (Preparation); PRP (Properties); USES (Uses) 1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE) L50 ANSWER 2 OF 58 REGISTRY COPYRIGHT 2005 ACS on STN **481774-46-3** REGISTRY Green fluorescent protein [65-glycine, 68-leucine, 72-alanine] (Aeguorea CN victoria) (9CI) (CA INDEX NAME) FS PROTEIN SEQUENCE SQL 238 SEO 1 MSKGEELFTG VVPILVELDG DVNGHKFSVS GEGEGDATYG KLTLKFICTT 51 GKLPVPWPTL VTTFGYGLQC FARYPDHMKR HDFFKSAMPE GYVOERTIFF 101 KDDGNYKTRA EVKFEGDTLV NRIELKGIDF KEDGNILGHK LEYNYNSHNV 151 YIMADKQKNG IKVNFKIRHN IEDGSVQLAD HYQQNTPIGD GPVLLPDNHY 201 LSTQSALSKD PNEKRDHMVL LEFVTAAGIT HGMDELYK --------HITS AT: 1-238 MF Unspecified

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CN
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L50 ANSWER 18 OF 58 REGISTRY COPYRIGHT 2005 ACS on STN
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L50 ANSWER 22 OF 58 REGISTRY COPYRIGHT 2005 ACS on STN
    481774-26-9 REGISTRY
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**RELATED SEQUENCES AVAILABLE WITH SEQLINK**
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L50 ANSWER 25 OF 58 REGISTRY COPYRIGHT 2005 ACS on STN
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L50 ANSWER 27 OF 58 REGISTRY COPYRIGHT 2005 ACS on STN
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L50 ANSWER 28 OF 58 REGISTRY COPYRIGHT 2005 ACS on STN
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L50 ANSWER 31 OF 58 REGISTRY COPYRIGHT 2005 ACS on STN
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1351	LFYEILCSLI	NPKRKDTRGF	SHFAEVTENF		GSPGEKSKTI
1401	LDSCPYLSIL	ALHWYPQQIN	GHKFEGKEGD		VQDAEIMAGK
1451	STCKLVQFTE	YSSQQWFISG	NNLPTLKNKV		QLLTNDNEVL
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1551	MSVYAVYART	DNLSSYNEAF	FTSGFICISG		CARYSMFAAK
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1651	IQSVNFWYVL	VMNDEHTERR	YLLFFLLSWG		VILKGIYHQS
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OTHER NAMES:
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   GenBank AAF94608 (Translated from: GenBank AE004223)
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L50 ANSWER 35 OF 58 REGISTRY COPYRIGHT 2005 ACS on STN

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OTHER NAMES:

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151 DLKQLPSQYQ YLSDTPADNA LVSKIQPLVV DILRQTANGM DEGEMKHALL

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   (9CI)
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       251 FVDREHTGIL IQIEPLKEIR KLILAFPMPS TESYYQKKPL SYFAHLIGYE
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       351 QSLFQTLNLI ATQGLQAWRY QEKRAVLESA FRFQETQRPL DMVSHLVVNM
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    451 OWYFTPYSVR PFTTEOLHRF ROPLDLPISL PEPNPFICYD LDPSEVKESH
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       601 LRKFAQRDFQ PKRFATIKQQ MTRNWRNAAH DKPISQLFNA MTGLLQPNNP
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       701 KDALRVQGQT YEESLRPLVM LGKSGTFORE VOCOODDSAI VVYYOSHEVS
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   N16961 gene VC1817) (9CI) (CA INDEX NAME)
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OTHER NAMES:
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   (9CI)
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OTHER NAMES:
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^{**}RELATED SEQUENCES AVAILABLE WITH SEQLINK**

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L50 ANSWER 46 OF 58 REGISTRY COPYRIGHT 2005 ACS on STN
RN
     290392-99-3 REGISTRY
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     Dehydrogenase, malate (Vibrio cholerae strain N16961 gene VC0432) (9CI)
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OTHER NAMES:
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    272763-00-5 REGISTRY
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   Protein JSAP1c (JNK/SAPK-associated protein-1c) (mouse) (9CI) (CA INDEX
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   12: PN: WO0031132 SEQID: 11 claimed protein
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   JSAP1b protein (mouse)
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   (9CI) (CA INDEX NAME)
OTHER NAMES:
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   GenBank AAF18575 (Translated from: GenBank AF115393)
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L50
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CN
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CN
   Protein (human clone 784CIF2B 161 precursor)
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SQL 479
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Source | Reference
Not Given | WO2001053312
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          2 REFERENCES IN FILE CAPLUS (1907 TO DATE)
   ANSWER 52 OF 58 REGISTRY COPYRIGHT 2005 ACS on STN
L50
   247916-06-9 REGISTRY
RN
CN
   Protein mAKAP (muscle A-kinase anchoring protein) (Rattus norvegicus
   strain Sprague-Dawley) (9CI) (CA INDEX NAME)
OTHER NAMES:
CN
   GenBank AAD39150
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   211809-45-9 REGISTRY
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L50 ANSWER 54 OF 58 REGISTRY COPYRIGHT 2005 ACS on STN
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L50 ANSWER 56 OF 58 REGISTRY COPYRIGHT 2005 ACS on STN
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  ANSWER 57 OF 58 REGISTRY COPYRIGHT 2005 ACS on STN
1.50
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   Phosphatase, adenosine tri- (Helicobacter pylori clone pBHpC8 gene copA)
   (9CI)
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          1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
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SEARCH CHARGES	0.00	229.20
DISPLAY CHARGES	0.00	435.82
OTHER CHARGES	. 0.00	10.50
FULL ESTIMATED COST	2.45	1044.75
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CA SUBSCRIBER PRICE	0.00	-9.49

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